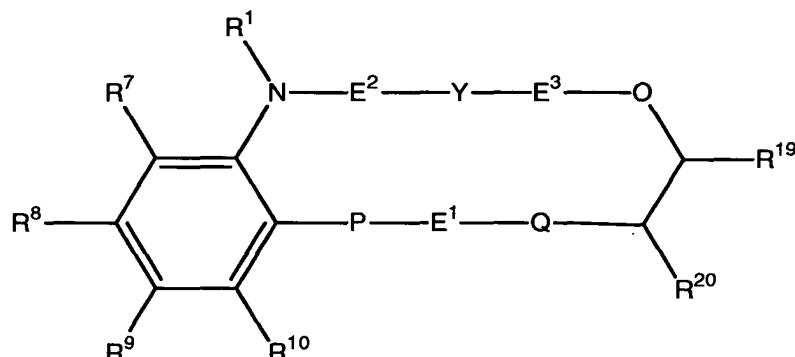


CLAIMS

WHAT IS CLAIMED IS:

5

1. A crown ether chelating compound having formula:



wherein

10 Y is O, S, NR⁴ or is absent, wherein R⁴ is selected from the group consisting of H, -L-R_x, -L-S_c, -L-DYE, C₁-C₁₈ alkyl, aryl and heteroaryl ring system, which alkyl or ring system is optionally substituted by halogen, azido, nitro, nitroso, amino, C₁-C₆ alkylamino, C₂-C₁₂ dialkylamino, cyano, -L-R_x, -L-S_c, -L-DYE, C₁-C₆ alkyl or C₁-C₆ alkoxy that is itself optionally substituted by halogen, amino, hydroxy, -(SO₂)-R¹⁵, -(SO₂)-O-R¹⁵, -(C=O)-R¹⁵, -(C=O)-O-R¹⁶,
15 or -(C=O)-NR¹⁷R¹⁸; wherein

R¹⁵ is selected from the group consisting of H, C₁-C₆ alkyl, -L-R_x, -L-S_c and -L-DYE;

R¹⁶ is selected from the group consisting of H, C₁-C₆ alkyl, benzyl, a biologically compatible esterifying group, a biologically compatible salt, -L-R_x, -L-S_c and -L-DYE;

20 R¹⁷ and R¹⁸ are independently selected from the group consisting of H, C₁-C₆ alkyl, C₁-C₆ carboxyalkyl, alpha-acyloxyalkyl, trialkylsilyl, a biologically compatible salt, -L-R_x, -L-S_c and -L-DYE; or R¹⁷ and R¹⁸ taken in combination form a 5- or 6-membered aliphatic ring that optionally incorporates an oxygen atom;

each L is independently a covalent linkage;

each R_x is independently a reactive group;

25 each S_c is independently a conjugated substance;

each DYE is independently a reporter molecule;

P and Q are independently O, S or NR³, wherein each R³ is independently H or C₁-C₆ alkyl;

E¹, E², and E³ are independently -(CR⁵)₂-, -(C(O)CH₂)_n-, -(CR⁵)_nO(CR⁵)_n- or E² is absent, where n = 2, 3 or 4, and each R⁵ is independently H or CH₃, or two R⁵ moieties on adjacent carbons of one or more of E¹, E² or E³, when taken in combination, form a 5- or 6-membered

5 aliphatic ring;

R¹ is selected from the group consisting of -L-R_X, -L-S_C, -L-DYE, C₁-C₁₈ alkyl and C₇-C₁₈ arylalkyl, each of which is optionally substituted by halogen, azido, nitro, nitroso, amino, hydroxy, cyano, C₁-C₆ alkoxy, an aryl or heteroaryl ring system, -(SO₂)-R¹⁵, -(SO₂)-O-R¹⁵, -(C=O)-R¹⁵, -(C=O)-O-R¹⁶, -(C=O)-NR¹⁷R¹⁸, C₁-C₆ alkylamino, C₂-C₁₂ dialkylamino, C₁-C₆

10 alkyl or C₁-C₆ alkoxy, each of which is itself optionally substituted by halogen, amino (-NR¹⁷R¹⁸), hydroxy, -(SO₂)-R¹⁵, -(SO₂)-O-R¹⁵, -(C=O)-R¹⁵, -(C=O)-O-R¹⁶ or -(C=O)-NR¹⁷R¹⁸;

R¹⁹ and R²⁰ are independently selected from the group consisting of H, halogen, azido, nitro, nitroso, amino, cyano, -L-R_X, -L-S_C, -L-DYE, C₁-C₆ alkyl and C₁-C₆ alkoxy, each of which is itself optionally substituted by halogen, amino, hydroxy, -(SO₂)-R¹⁵, -(SO₂)-O-R¹⁵, -(C=O)-

15 R¹⁵, -(C=O)-O-R¹⁶, or -(C=O)-NR¹⁷R¹⁸;

or R¹⁹ and R²⁰ taken in combination form a fused six-membered benzo moiety that is optionally substituted by halogen, azido, nitro, nitroso, amino, cyano, -L-R_X, -L-S_C, -L-DYE, C₁-C₆ alkyl or C₁-C₆ alkoxy, each of which is itself optionally substituted by halogen, amino, hydroxy, -(SO₂)-R¹⁵, -(SO₂)-O-R¹⁵, -(C=O)-R¹⁵, -(C=O)-O-R¹⁶, or -(C=O)-NR¹⁷R¹⁸;

20 R⁷, R⁸, R⁹ and R¹⁰ are independently selected from the group consisting of H, halogen, azido, nitro, nitroso, amino, cyano, -L-R_X, -L-S_C, -L-DYE, C₁-C₆ alkyl or C₁-C₆ alkoxy, each of which is itself optionally substituted by halogen, amino, hydroxy, -(SO₂)-R¹⁵, -(SO₂)-O-R¹⁵, -(C=O)-R¹⁵, -(C=O)-O-R¹⁶, or -(C=O)-NR¹⁷R¹⁸;

or any two adjacent substituents R⁷-R¹⁰, taken in combination, form a fused six-membered

25 benzo moiety, which is optionally substituted by halogen, azido, nitro, nitroso, amino, cyano, -L-R_X, -L-S_C, -L-DYE, C₁-C₆ alkyl or C₁-C₆ alkoxy, each of which is optionally substituted by halogen, amino, hydroxy, -(C=O)-R¹⁵, -(C=O)-O-R¹⁶, or -(C=O)-NR¹⁷R¹⁸;

or any two adjacent substituents R⁷-R¹⁰, or R¹⁹ and R²⁰, taken in combination with each other, form a fused DYE.

30 2. The compound according to Claim 1, wherein said P, Q and Y are O.

3. The compound according to Claim 2, wherein said E¹, E², and E³ are each -(CR⁵)_n wherein R⁵ is H and each n is 2.
4. The compound according to Claim 3, wherein at least one of said R¹, R⁷, R⁸, R⁹, R¹⁰, R¹⁹ or R²⁰ is -L-Rx, -L-Sc or -L-DYE or R⁸ in combination with R⁹ form a fused DYE.
5
5. The compound according to Claim 4, wherein said L is a single covalent bond, or a covalent linkage that is linear or branched, cyclic or heterocyclic, saturated or unsaturated, having 1-20 nonhydrogen atoms selected from the group consisting of C, N, P, O and S; and are composed of any combination of ether, thioether, amine, ester, carboxamide, sulfonamide, hydrazide bonds and aromatic or heteroaromatic bonds.
10
6. The compound according to Claim 4, wherein said -Rx is selected from the group consisting of an acrylamide, an activated ester of a carboxylic acid, a carboxylic ester, an acyl azide, an acyl nitrile, an aldehyde, an alkyl halide, an anhydride, an aniline, an amine, an aryl halide, an azide, an aziridine, a boronate, a diazoalkane, a haloacetamide, a halotriazine, a hydrazine, an imido ester, an isocyanate, an isothiocyanate, a maleimide, a phosphoramidite, a reactive platinum complex, a silyl halide, a sulfonyl halide, a thiol and a photoactivatable group.
15
7. The compound according to Claim 6, wherein said -Rx is selected from the group consisting of carboxylic acid, succinimidyl ester of a carboxylic acid, hydrazide, amine and a maleimide.
20
8. The compound according to Claim 4, wherein said -Sc is selected from the group consisting of an amino acid, a peptide, a protein, a polysaccharide, a nucleoside, a nucleotide, an oligonucleotide, a nucleic acid, a hapten, a psoralen, a drug, a hormone, a lipid, a lipid assembly, a synthetic polymer, a polymeric microparticle, a biological cell or a virus.
25
9. The compound according to Claim 8, wherein said -Sc is selected from the group consisting of an antibody or fragment thereof, an avidin or streptavidin, a biotin, a blood component protein, a dextran, an enzyme, an enzyme inhibitor, a hormone, an IgG binding protein, a fluorescent protein, a growth factor, a lectin, a lipopolysaccharide, a microorganism, a metal binding protein, a metal chelating
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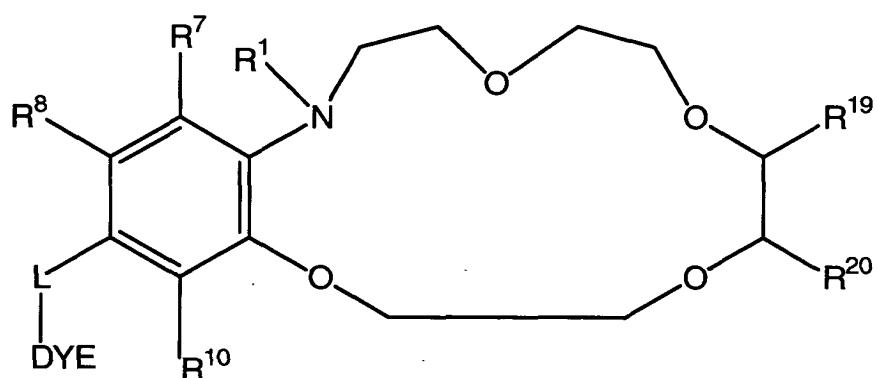
moiety, a non-biological microparticle, a peptide toxin, a phosphatidylserine-binding protein, a structural protein, a small-molecule drug, or a tyramide.

10. The compound according to Claim 5, wherein said -DYE is selected from the group
5 consisting of xanthene, boropolyazaindacene, carbocyanine, benzofuran,
quinazolinone, indole, a benzazole, oxazine, and coumarin.

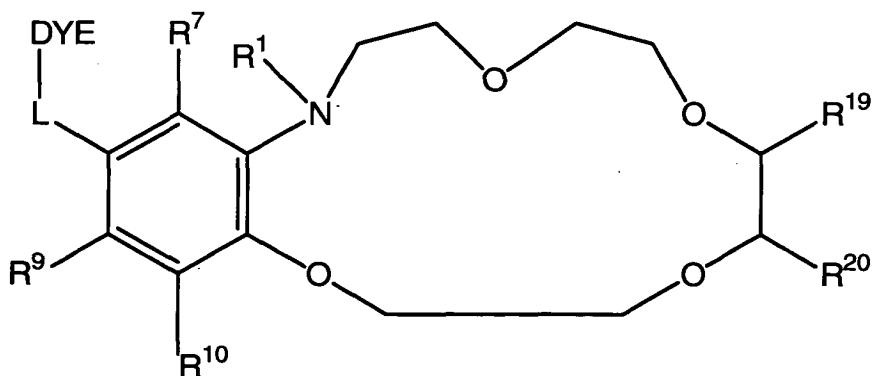
11. The compound according to Claim 10, wherein said -DYE moiety is independently
10 substituted by a lipophilic group.

12. The compound according to Claim 11, wherein said lipophilic group is an AM or
acetate ester.

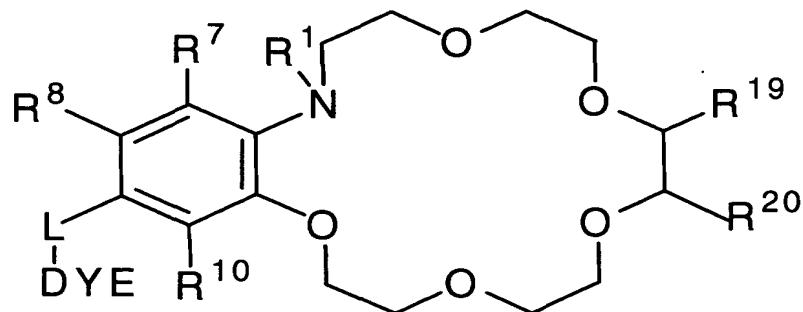
13. The compound according to Claim 1, wherein said compound is selected from the
15 group consisting of



Formula (II)(a),

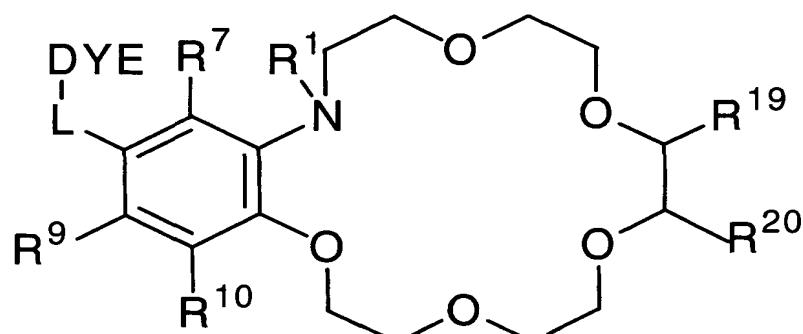


Formula (II)(b),

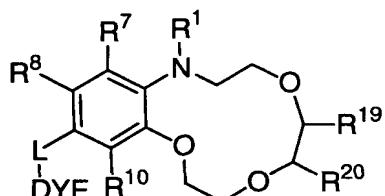


Formula (II)(c),

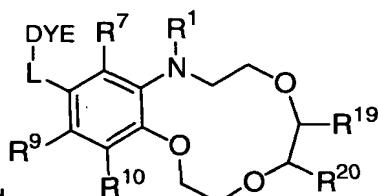
5



Formula (II)(d),



Formula (II)(e) and



Formula (II)(f).

10

14. The compound according to Claim 13, wherein said DYE is selected from the group consisting of boropolyazaindacene, xanthene and indole.

15 15. The compound according to Claim 13, wherein said DYE moiety is independently substituted by a lipophilic group.

16. The compound according to Claim 15, wherein said lipophilic group is an AM or acetate ester.

20

17. The compound according to Claim 13, wherein R⁷, R⁸, R⁹, R¹⁰, R¹⁹ and R²⁰, when present, are H.

18. The compound according to Claim 17, wherein R¹ is C₁-C₆ alkyl that is substituted one or more times by amino (-NR¹⁷R¹⁸), -(C=O)-O-R¹⁶ or -(C=O)-NR¹⁷R¹⁸.

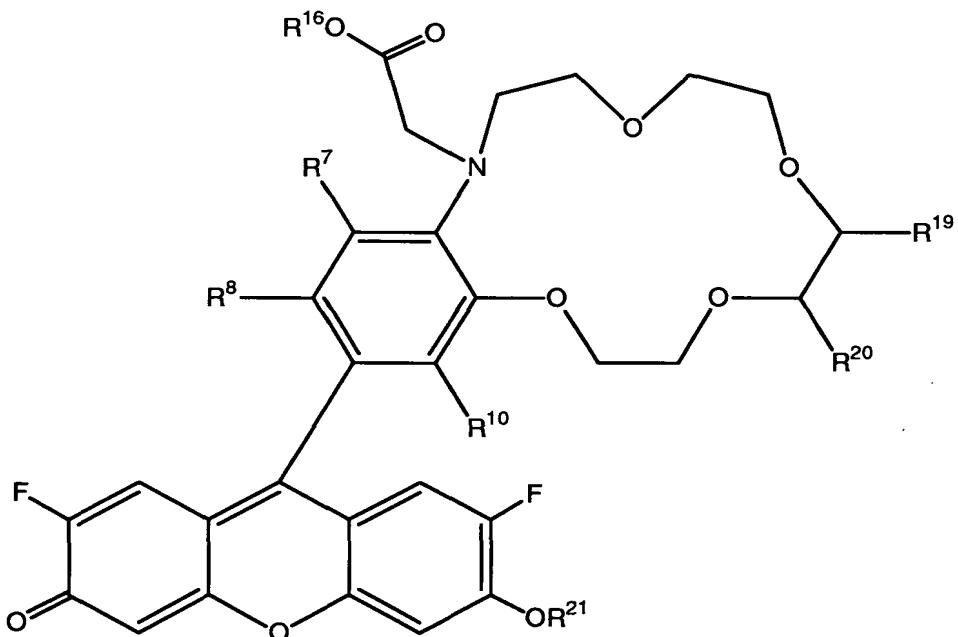
5 19. The compound according to Claim 18, wherein said R¹ is methyl or ethyl.

20. The compound according to Claim 19 wherein said R¹⁶ is selected from the group consisting of H, C₁-C₆ alkyl, benzyl, a biologically compatible esterifying group, and a biologically compatible salt.

10 21. The compound according to Claim 20 wherein said R¹⁶ is methyl.

15 22. The compound according to Claim 18 wherein said R¹⁷ and R¹⁸ are each methyl.

23. A compound having formula:



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wherein R¹⁶ is selected from the group consisting of H, C₁-C₆ alkyl, benzyl, a biologically compatible esterifying group, and a biologically compatible salt;

R¹⁹ and R²⁰ are selected from the group consisting of H, halogen, azido, nitro, nitroso, amino, cyano, -L-R_x, -L-S_C, -L-DYE, C₁-C₆ alkyl and C₁-C₆ alkoxy, each of which is itself optionally substituted by halogen, amino, hydroxy, -(SO₂)-R¹⁵, -(SO₂)-O-R¹⁵, -(C=O)-R¹⁵, -(C=O)-O-R¹⁶ and -(C=O)-NR¹⁷R¹⁸;

or R¹⁹ and R²⁰ taken in combination form a fused six-membered benzo moiety that is optionally substituted by halogen, azido, nitro, nitroso, amino, cyano, -L-R_x, -L-S_C, -L-DYE, C₁-C₆ alkyl or C₁-C₆ alkoxy, each of which is itself optionally substituted by halogen, amino, hydroxy, -(SO₂)-R¹⁵, -(SO₂)-O-R¹⁵, -(C=O)-R¹⁵, -(C=O)-O-R¹⁶, or -(C=O)-NR¹⁷R¹⁸;

R¹⁵ is selected from the group consisting of H, C₁-C₆ alkyl, -L-R_x, -L-S_C and -L-DYE;

R¹⁶ is selected from the group consisting of H, C₁-C₆ alkyl, benzyl, a biologically compatible esterifying group, a biologically compatible salt, -L-R_x, -L-S_C and -L-DYE;

R¹⁷ and R¹⁸ are independently selected from the group consisting of H, C₁-C₆ alkyl, C₁-C₆ carboxyalkyl, alpha-acyloxyalkyl, trialkylsilyl, a biologically compatible salt, -L-R_x, -L-S_C and -L-DYE; or R¹⁷ and R¹⁸ taken in combination form a 5- or 6-membered aliphatic ring that optionally incorporates an oxygen atom;

each L is independently a covalent linkage;

each R_x is independently a reactive group;

each S_C is independently a conjugated substance;

each DYE is independently a reporter molecule;

R⁷, R⁸, and R¹⁰ are independently selected from the group consisting of H, halogen, azido, nitro, nitroso, amino, cyano, -L-R_x, -L-S_C, -L-DYE, C₁-C₆ alkyl and C₁-C₆ alkoxy, each of which is optionally substituted by halogen, amino, hydroxy, -(SO₂)-R¹⁵, -(SO₂)-O-R¹⁵, -(C=O)-R¹⁵, -(C=O)-O-R¹⁶, or -(C=O)-NR¹⁷R¹⁸;

or R⁷ taken in combination with R⁸ form a fused six-membered benzo moiety, which is optionally substituted by halogen, azido, nitro, nitroso, amino, cyano, -L-R_x, -L-S_C, -L-DYE, C₁-C₆ alkyl or C₁-C₆ alkoxy, each of which is optionally substituted by halogen, amino, hydroxy, -(C=O)-R¹⁵, -(C=O)-O-R¹⁶, or -(C=O)-NR¹⁷R¹⁸; and,

R^{21} is selected from the group consisting of H, C₁-C₁₈ alkyl, C₇-C₁₈ arylalkyl and a lipophilic group each alkyl is optionally substituted by -(C=O)-R¹⁵, -(C=O)-O-R¹⁶, or C₁-C₆ alkoxy.

24. The compound according to Claim 23, wherein said R⁷, R⁸, and R¹⁰ are H.
5
25. The compound according to Claim 24, wherein said R¹⁹ and R²⁰ are H.
10
26. The compound according to Claim 25, wherein said R¹⁶ is methyl or a biologically compatible esterifying group.
15
27.. The compound according to Claim 23 wherein said R¹⁶, R¹⁹ or R²⁰ is -L-Sc.
28. A composition comprising:
15
a. a compound according to any one of Claims 1-27; and,
b. a metal ion that is capable of being chelated by said compound.
29. The composition according to Claim 28, wherein said metal ion is selected from the group consisting of Na⁺, Li⁺, K⁺, Ca⁺, Zn⁺ and Rb⁺.
20
30. A method for binding a target metal ion in a sample, comprising steps of:
a. contacting said sample with a metal chelating compound according to any one of Claims 1-27
b. incubating said sample and said metal chelating compound for sufficient time to allow said compound to chelate said target metal ion whereby said metal ion is bound.
25
31. The method according to Claim 30, wherein said method further comprises illuminating said metal chelating compound with a suitable light source whereby said target ion is detected with the proviso that at least one of R¹, R⁴, R⁷, R⁸, R⁹, R¹⁰, R¹⁹ or R²⁰ is -L-DYE or at least two of R⁷-R¹⁰ or R¹⁹ and R²⁰, taken in combination, form a fused DYE.
30
32. The method according to Claim 31, wherein said target metal ion is selected from the group consisting of Na⁺, Li⁺, K⁺, Ca⁺, Zn⁺ and Rb⁺.
35

33. The method according to Claim 32, wherein said target metal ion is Na^+ .

34. The method according to Claim 32, wherein said sample comprises living cells, cellular components, proteins, peptides, buffer solutions or biological fluids.

5

35. A method for binding and detecting target ions in a live cell, said method comprises:

- a) contacting a sample of live cells with a crown ether compound according to any one of Claims 1-27 with the proviso that said compound comprise a DYE moiety and at least one lipophilic group;
- 10 b) incubating said sample and said crown ether chelate compound for sufficient time to allow said compound to chelate said target metal ion; and,
- c) illuminate said sample with an appropriate wavelength whereby said target ion is detected in a live cell.

15 36. The method according to Claim 35, wherein said DYE moiety is substituted by a lipophilic group.

37. The method according to Claim 36 wherein said lipophilic group is an AM or acetate ester.

20 38. A kit for binding a metal ion in a sample, comprising:
a compound according to any one of Claims 1-27; and, comprising one or more components selected from the group consisting of a calibration standard of a metal ion, an ionophore, a fluorescent standard, an aqueous buffer solution and an organic solvent.

25